and by adding in its place ''Granary Chambers, 37–39 Burton St., Melton Mowbray, Leicestershire LE13 1AF, England''.

Dated: October 29, 1996.

Andrew J. Beaulieau,

Deputy Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 96–29388 Filed 11–15–96; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Parts 510 and 558

Animal Drugs, Feeds, and Related Products; Tylosin

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to remove that portion reflecting approval of a new animal drug application (NADA) held by Countrymark Cooperative, Inc. (formerly Indiana Farm Bureau Cooperative Association, Inc.). The NADA provides for use of a tylosin Type A medicated article for making a tylosin Type C medicated swine feed. In a notice published elsewhere in this issue of the Federal Register, FDA is withdrawing approval of the NADA.

EFFECTIVE DATE: November 29, 1996.

FOR FURTHER INFORMATION CONTACT: Mohammad I. Sharar, Center for Veterinary Medicine (HFV–216), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1722.

SUPPLEMENTARY INFORMATION: In a notice published elsewhere in this issue of the Federal Register, FDA is withdrawing approval of NADA 125–226 held by Countrymark Cooperative, Inc., 950 North Meridian St., Indianapolis, IN 46204–3909 (formerly Indiana Farm Bureau Cooperative Association, Inc., 120 East Market St., Indianapolis, IN 46204). The NADA provides for use of tylosin Type A medicated articles to make tylosin Type C medicated swine feeds. Countrymark Cooperative, Inc., voluntarily requested withdrawal of approval of the NADA because it no longer makes Type A medicated articles for use in medicated feeds. This document removes the entry in 21 CFR 558.625(b) to reflect the withdrawal of approval of this NADA.

This NADA was originally held by Indiana Farm Bureau Cooperative Association, Inc. The regulations had not been amended in § 510.600(c) (21 CFR 510.600(c)) to reflect the sponsor change to Countrymark Cooperative. At this time, Indiana Farm Bureau Cooperative Association is no longer the sponsor of any approved NADA's. Therefore, § 510.600(c) is amended to remove the entries for the firm.

List of Subjects

21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

21 CFR Part 558

Animal drugs, Animal feeds.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR parts 510 and 558 are amended as follows:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: Secs. 201, 301, 501, 502, 503, 512, 701, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e).

§ 510.600 [Amended]

2. Section 510.600 *Names, addresses, and drug labeler codes of sponsors of approved applications* is amended in paragraph (c)(1) by removing the entry for ''Indiana Farm Bureau Cooperative Association, Inc.,'' and in paragraph (c)(2) by removing the entry for ''021502.''

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

3. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

§ 558.625 [Amended]

4. Section 558.625 *Tylosin* is amended by removing and reserving paragraph (b)(76).

Dated: October 18, 1996.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 96–29389 Filed 11–15–96; 8:45 am]

BILLING CODE 4160–01–F

Health Care Financing Administration

42 CFR Part 413

[BPD–805–CN]

RIN 0938–AG68

Medicare and Medicaid Programs; New Payment Methodology for Routine Extended Care Services Provided in a Swing-Bed Hospital; Correction

AGENCY: Health Care Financing Administration (HCFA), HHS.

ACTION: Correction notice.

SUMMARY: This document corrects the final rule published October 3, 1996 (61 FR 51611) that revised the methodology for payment of routine extended care services furnished in a swing-bed hospital. The final rule also revised the regulations concerning the method used to allocate hospital general routine inpatient service costs for purposes of determining payments to swing-bed hospitals.

EFFECTIVE DATE: These corrections are effective as of November 4, 1996.

FOR FURTHER INFORMATION CONTACT: John Davis, (410) 786–0008.

SUPPLEMENTARY INFORMATION: We are making the following corrections to the October 3, 1996 final rule (61 FR 51611):

1. On page 51612, in the first column, fourth line from the bottom, the duplicate word ''harmless'' is deleted.

2. On page 51612, in the third column, lines 16 and 17, the phrase ''ending on or after June 30, 1989 and through May 31, 1990'' is corrected to read ''ending on or after June 30, 1989 through May 31, 1990''.

3. On page 51615, in the third column, lines 30 and 31, the phrase ''we are changing to the out method'' is corrected to read ''we are changing to the carve-out method''.

4. On page 51616, in the first column, under Subpart D, item number 2, the amendatory language is corrected by adding the phrase ''the introductory text of paragraph (a)(1)(ii);'' after the phrase ''Section 413.53 is amended by revising''.

(Catalog of Federal Domestic Assistance Program No. 93.773 Medicare—Hospital Insurance Program; No. 93.778, Medical Assistance Program)

Dated: November 7, 1996.

Neil J. Stillman,

Deputy Assistant Secretary for Information Resources Management.

[FR Doc. 96–29398 Filed 11–15–96; 8:45 am]

BILLING CODE 4120–01–P